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Physical and Biological Mechanisms Influencing the Development and Evolution of Sedimentary Structure



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INTRODUCTION

This document is the final summary report for research conducted under Naval Research Laboratory (NRL) parent grant N00014-93-1-6004, "Physical and biological mechanisms influencing the development and evolution of sedimentary structure," funded through the Coastal Benthic Boundary Layer Special Research Program during fiscal years 1993 and 1994.

We participated in three research cruises in winter and spring 1993, and one in summer 1994; analyses were carried out through 1993 and 1994, and results presented in 1994 and 1995. The description of methods, results, and discussion in this report reflects that chronology.

GENERAL SUMMARY OF WORK DURING THE FIRST YEAR

Research work completed during 1993 included the collection of samples in Eckernfoerde Bucht and the beginning of laboratory analyses for those samples. During March-May 1993, we participated in three cruises. The sample sites were concentrated at the deployment locations of acoustic and boundary-layer towers, as well as positions farther northeastward and southwestward within the bay (Fig. 1). Replicate cores (4-7) were collected at 14 stations, and many of the stations were reoccupied on different cruises (Table 1). Some of our sample processing was completed in the field, but most awaited the return of samples to our laboratory.

They were returned to the US in August, and analyses of sedimentological, radiochemical and biological properties were begun.

Our work has been coordinated with the dynamical boundary-layer measurements of our co-PI, Don Wright and his research group at VIMS, as well as the acoustical observations of Darrell Jackson and his research group at University of Washington. The combined results are providing an understanding of physical and biological reworking for the seabed in Eckernfoerde Bay, and how this has affected sedimentological characteristics and the resultant acoustical properties.

Sedimentary structure in Eckernfoerde Bay is generally characterized by physical laminations, although biogenic mottling is present and becomes more common northeastward (toward the mouth). The benthos consist of an opportunistic community dominated by spionid polychaetes and tellinid bivalves, both surface deposit feeders. Profiles of ²³⁴Th demonstrate that the community mixing depth is 1 cm or less. This observation combined with a relatively rapid accumulation rate (>3 mm y⁻¹) allows many physical laminae to pass through the surface mixed layer intact, and to be preserved in the underlying portions of the seabed.

Description of Coring Stations

Most of the cores were collected at locations where the APL tower and the VIMS tetrapod were deployed, as shown below (Figure 1, Table 1).

| Table 1. Station locations. |
|-----------------------------|
|-----------------------------|

| | <u>BS-1</u> | <u>BS-2</u> | <u>BS-3</u> |
|--------------------------------------|-------------------------|-------------------|-----------------|
| APL/VIMS deployment site March-April | stations A,B,C,D,E,F | stations D,F | station D |
| APL/VIMS deployment site April-May | none | stations N,O,P | stations N,O |

Cruise Number

Two locations near Mittelgrund contained glacial sand and pebbles, and had a more diverse benthic community. The location at the west end of Eckernfoerde (near the Navy Base) had a large contribution of transported debris (sand, twigs, leaves, eel grass).

Dist Special

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Cores were examined by observations of X-radiographs, microfabric, grain size, 234 Th , and 210 Pb. Slabs for X-radiography were obtained by pushing three-sided acrylic trays (2.5 x 15 cm cross-section) into core boxes on deck, then inserting the fourth side, so as to minimize fabric distortion (Kuehl and others, 1985). Slabs were fixed with formalin, sealed, and X-rayed aboard ship. Microfabric subsamples were obtained from X-ray slabs, and prepared using resin impregnation and petrographic thin-sectioning (Kuehl and others, 1988). Subsamples for grain-size measurement were disaggregated with 30% H_2O_2 (also removing organic matter; Ingram, 1971). Granulometric measurements were made with a Micromeritics ET-5100 Sedigraph following the methods of Coakley and Syvitski (1991), and were cross-checked with standard sieve, pipette, and settling-column techniques. 234 Th activities ($t_{1/2}$ = 24 days) in the upper 10 cm were determined by γ -spectroscopy (63 keV peak: Buesseler and others, 1992). 210 Pb activities were measured by wet chemical leaching of samples to extract the granddaughter 210 Po, which was then analyzed by α - spectroscopy (Nittrouer and others, 1979). Excess activities of 210 Pb and 234 Th were determined by comparison with supported background activities.

Benthic Biological Sampling

The benthos subsample cores were divided into 0-2 and 2-10 cm layers, and sieved through a 500 µm sieve. Animals collected on the sieve were then preserved in glass jars with 4% seawater formaldehyde and rose bengal. Animals were counted and identified to the lowest taxonomic unit.

A bioturbation experiment was established to compare vertical and horizontal particle bioturbation rates in a variety of sediments and faunal communities. Fifteen-cm diameter cores (see Table 2) were brought to shore and placed in a running seawater system. A 50-ml suspension of a second color of fluorescent particles (in seawater) was added to each core. These particles settled overnight onto the sediment surface.

Control cores were broken down during the following two days. There was one control core from each station. Nine to 12 subcores (small syringes) were taken from each control core. Each subcore was vertically sectioned at 7 depth intervals of 0-0.5, 0.5-1, 1-1.5, 1.5-2, 2-3, 3-4,

and 4-5 cm. Cores for porosity also were taken and divided in the same manner. Small x-ray slabs were obtained from each core.

Table 2. Description of samples for bioturbation experiment

| Date sampled | Station | Box core # | Comments | |
|--------------|---------|------------|-------------------|------------------|
| 27 April 93 | Ι | 106, A, B | 4 small box cores | Transition Site |
| | | 107, C, D | | |
| 28 April 93 | D | 119 A-C | Large box core | Old Tower Site |
| 28 April 93 | H | 114 A-D | Large box core | Navy Base Site |
| 28 April 93 | N | 128 A-D | Large box core | New Tower Site |
| 29 April 93 | Ο | 143 A-C | Large box core | New Tower Site |
| 29 April 93 | G | 146 A, B | Small box core | Mittelgrund Site |
| | | 147 C | | |
| 29 April 93 | F | 151 A-D | Large box core | Old Tower Site |

Table 3. Station locations and core subsamples, BS4.

| Station | Sample Type | Date/Time | Location ¹ | <u>Latitude N</u> | <u>LongitudeE</u> | Water Depth | Subsampling ² |
|---------|--------------|----------------|-----------------------|-------------------|-------------------|-------------|--------------------------|
| 602 | Box core | 06/27/94 14:45 | N | 54 23 43.9 | 09 59 31.0 | 25 m | XR, Pb, BIO |
| 604 | Gravity core | 06/27/94 18:15 | N | 54 29 44.9 | 09 59 29.4 | 25 m | Pb |
| 606 | Gravity core | 06/28/94 09:47 | F | 54 29 25,54 | 09 58 59.05 | 25.5 m | Pb |
| 607 | Gravity core | 06/28/94 10:16 | F | 54 29 22.97 | 09 59 00.03 | 25.5 | Pb |
| 611 | Box core | 06/28/94 11:22 | F | 54 29 24.4 | 09 59 01.2 | 25 m | Pb, XR, C, BIO |
| 620 | Box core | 06/29/94 10:41 | HG | 54 32 13.26 | 10 03 33.50 | 27 m | Pb, XR, BIO, C |
| 621 | Gravity core | 06/29/94 11:16 | HG | 54 32 12.0 | 10 03 36.0 | 28 m | C, Pb |
| 623 | Gravity core | 06/29/94 12:05 | HG | 54 32 12.3 | 10 03 29.1 | 28 m | C, Pb |
| 630 | Box core | 06/30/94 08:31 | TR | 54 30 00.95 | 09 58 33.69 | 25 m | XR, BIO |
| 631 | Box core | 06/30/94 08:51 | TR | 54 30 10.60 | 09 58 31.87 | 21 m | XR, BIO |
| 633 | Box core | 06/30/94 09:11 | TR | 54 30 12.50 | 09 58 35.83 | 21 m | XR, BIO |
| 635 | Box core | 07/01/94 09:17 | TR | 54 30 06.16 | 09 58 24.68 | 22 m | XR, BIO |
| 636 | Box core | 07/01/94 09:32 | TR | 54 30 07.59 | 09 58 15.53 | 17 m | XR, BIO |
| 637 | Box core | 07/01/94 09:53 | TR | 54 30 03.66 | 09 58 00.41 | 14 m | XR, BIO |
| 648 | Box core | 07/02/94 am | F? | 54 29 26.15 | 09 58 14.00 | 25 m | Th |

Salinity and temperature were monitored during the course of the experiment.

Experimental cores were incubated in the running seawater system for approximately 2 weeks, then were broken down in the same manner as the controls. After subcores were taken from the experimental cores, the remaining sediment (~70% of the total) was sieved through a 500-µm

 $^{^{1}}$ N,F = APL tower sites; HG = Hausgarten; TR = transect stations. 2 XR = X-ray slabs, also for microstructure; Pb = 210 Pb subsampling; C = 14 C subsamples; BIO = fecal pellet and/or organism subsamples; Th = 234 Th subsampling.

sieve, and the animals collected were preserved for later enumeration. Fluorescence from sediment subsamples was extracted and measured by fluorometry.

SEDIMENTOLOGY AND RADIOCHEMISTRY

Summary of work

Sediment in the central basin of Eckernfoerde Bay is characterized by beds of clay-sized material, with slightly coarser laminations (~10% silt and sand). Excess ²³⁴Th activity indicates that mixing is limited to the upper 8 mm of the seabed. Accumulation rates, measured by ²¹⁰Pb geochronology, range from 3 to over 11 mm yr⁻¹. The relatively high accumulation rates and absence of a significant mixed layer in the central bay allow fluctuations of sedimentological processes to be recorded in fine detail. Sediment-transport studies indicate that the laminated bedding is caused by alternating deposition of storm-suspended sediments from adjacent shallowwater areas and fair-weather supply of fine suspended material advected from the open Baltic.

X-radiography

X-radiographs of cores from the central basin of Eckernfoerde Bay show laminations in the upper 20-30 cm of the sea floor. The laminations are interbedded with thicker beds (several cm thick). Contacts between beds and laminations are diffuse. Laminations are generally more absorptive of X-rays than the thicker beds (thus appearing dark in positive prints: Fig. 2), and may be laterally discontinuous.

Microfabric

Two general types of microfabric were observed in cores from the central basin: pelletal fabrics (pellets > 10% of volume; percentages are approximate) and non-pelletal fabrics (pellets < 10% of volume). Of these, pelletal fabrics are by far the most common, and may be either matrix supported or pellet supported. Transitions from non-pelletal layers to overlying pelletal layers tend to be gradational, whereas contacts between non-pelletal and underlying pelletal layers are sharp (Fig. 3).

Pelletal fabrics tend to be anisotropic. Non-pelletal laminations sometimes display graded bedding (Fig. 3); other laminations may show no obvious grain-size variations between layers. In general, microstructural laminations correspond to laminations observed in X-radiographs, and pelletal fabrics in thin section are associated with thicker, homogenized beds seen in X-radiographs.

Grain-size distribution

Grain-size distributions tend to be polymodal, with the primary mode centered between 9.5 and 11.5 φ and a secondary mode, less than 10% by mass, centered between 3 and 6 φ (Fig. 4). Subtle coarsening in the median grain size of each mode is commonly associated with laminations observed in X-radiographs (Fig. 2), and non-pelletal, graded bedding in microfabric (Fig. 3). The coarsening associated with these laminations is accompanied by a 2-4% increase in the mass percentage of particles in the 3-6 φ size range.

Radiochemistry

Sediment accumulation rates in the central basin determined by ²¹⁰Pb geochronology range from 3 to over 11 mm per year (Fig. 5). Excess ²¹⁰Pb is present in the seabed to depths of 30-40 cm (Fig 5). Excess ²³⁴Th is restricted to the upper 5 mm of sediment, indicating that mixing is restricted to this thin veneer in the sea floor. No significant seasonal variability in sediment mixing depths was observed over winter, spring and summer cruises, although intensity may of mixing may change. Lateral variability of accumulation rates within the central basin was minimal.

Shallow penetration of sediment mixing results in the preservation of a detailed environmental record (cm-scale) in central-basin sediments. Long-term variations in ²¹⁰Pb profiles were observed in central-basin cores (Fig. 5), indicating historical shifts in either ²¹⁰Pb flux in coastal waters or changing sediment supply.

Sedimentary Event Layers

Once a sediment layer is deposited, it is then subject to modification by biological and physical mixing. In the absence of physical mixing, the degree of event-layer preservation in the

historical sedimentary record is a function of the deposited-layer thickness, the sediment accumulation rate, and the depth and rate of biological mixing (Guinasso and Schink, 1975; Nittrouer and Sternberg, 1981; Wheatcroft, 1990, 1994). Events layers thicker than the mixed layer can be generated by unusual events (from large storms or floods), and can overwhelm the benthos, interrupting biological mixing (Wheatcroft, 1994). (Biological mixing may be interrupted by other types of disturbances as well, such as anoxia, although the stratigraphic signature of temporary anoxia might be less obvious than a storm deposit.) The physical record of such an event would be a bed or lamination with a sharp lower boundary (resting on top of a bioturbated bed), a physically stratified lower layer, and a gradual increase upward in the intensity of bioturbation. This sequence has been widely recognized in coastal depositional environments (Frey and Howard, 1986; Leithold, 1989; Aigner and Reineck, 1982; Wheatcroft, 1994). The conceptual model is consistent with sedimentation in the central basin of Eckernfoerde Bay.

Depositional and Post-depositional Processes: Preserved Fabric

In order to assess the relationship between pulsed sediment deposition and rate of bioturbation, laminations were examined in thin section for evidence of progressive bioturbation with age. The laminations clearly preserved in thin sections (Fig. 3) have sharp lower contacts, and grade upward into progressively more pelletized sediment. This trend indicates that degree of pelletization can be used as an estimate of bioturbation intensity following event-layer deposition, when the pellets under study are resistant to breakdown. The thicknesses of unburrowed sediment in event layers are on the order of 3-5 mm, comparable to progressively pelletized layers. Each preserved lamination must have been initially thicker than the sediment mixing depth in the central basin, otherwise the lower boundary would not have been preserved.

Excess 234 Th activities (Fig. 5) were used to estimate the biodiffusion coefficient D_b (~ 0.3 cm² yr⁻¹) in central basin sediments, using a solution to the steady-state advection/diffusion equation (Aller and Cochran, 1976),

$$D_b = \lambda (z/\ln(C_0/C_z))^2$$

where λ is the ²³⁴Th decay constant (10.5 yr⁻¹), z is depth in the seabed (cm), C_0 is excess activity at the surface (decays per minute per gram of sediment: dpm/g), and C_z is excess activity at depth z (dpm/g). The thin surface mixed layer resulted in only two subsamples containing excess ²³⁴Th

(Fig. 5). Because of the rapid mixing rates relative to accumulation rates, biological mixing is assumed to control the penetration of excess ²³⁴Th into the seabed.

A method useful for assessing appropriate length and time scales of bioturbation is the decomposition of the biodiffusion coefficient into a mean step length and rest period (Wheatcroft and others, 1990) using the form $D_b = \delta^2/2\Omega$, where δ is mean step length, and Ω is mean rest period (yr). Assuming step lengths of 2 mm (appropriate to the benthic community, based on body length), the rest period is 24 days. To advect a 1 mm layer through the 5 mm mixed layer in less time than the rest period (and so preserve the layer) would require rapid deposition at rates equivalent to 7.5 cm yr⁻¹, one order of magnitude higher than the mean accumulation rates in the central basin (~7 mm yr⁻¹). Event layers exceeding the mixed-layer thickness satisfy these conditions.

Decadal-Scale Stratigraphic Record

Evidence of fluctuating accumulation rate is found on decadal scales, determined by ²¹⁰Pb geochronology (Fig. 5), as well as scales of single events, represented by non-pelletized laminations seen in thin sections (Fig. 4). Such fluctuations have been documented regionally (Balzer and others, 1986), and have been linked to winter-storm frequency and resultant resuspension and deposition (Khandriche et al, 1986). Shear velocities in the central basin have upper limits below critical erosion values (Friedrichs and Wright, in press), precluding significant physical sediment mixing. Biological mixing is seasonally intense, but is limited to depths (~8 mm) on the same order as annual accumulation rates (3-11 mm yr⁻¹). Thus, upper sediments of the central basin preserve a relatively high-resolution historical record (cm-scale) of sedimentary processes.

Conclusions

The sedimentary input to the central basin is controlled by pulsed supply of fine particles from both proximal and distal sources. Time-series benthic-boundary-layer observations (Friedrichs and Wright, in press) indicate physical sediment mixing to be insignificant. Biological mixing is intense, but presently limited to the upper 8 mm of seabed. Sediment pulses (storm layers) exceeding the mixed-layer thickness may be preserved as individual laminations. Slower

deposition results in intensely reworked and pelletized sediment (pelletal fabric). Particle and chemical tracers reaching the seafloor may be vertically resolved to increments of ~5 mm, preserving a detailed stratigraphic record.

Fluctuations in the frequency of sediment pulses cause variations in ²¹⁰Pb accumulation rates (3-11 mm yr⁻¹). Similarly, sedimentary fabric (mm and cm scales) displays subtle gradients in grain size and bioturbation (including pelletization) that correlate to fluctuations in sediment supply rate.

BENTHIC BIOLOGY

Summary of work

The benthic macrofaunal community of Eckernfoerde Bay was characterized in terms of abundance and community structure. The community was dominated by small surface deposit-feeding animals. Faunal abundances in spring 1993 were 30,000-80,000 m⁻², and were 2,000-9,000 m⁻² in summer 1994. Particle bioturbation was limited to the top 0.5-1.0 cm throughout Eckernfoerde Bay. Functional group classification, faunal abundances, organism size, and particle bioturbation are consistent with the hypothesis that the benthic community of Eckernfoerde Bay is controlled by a regular disturbance which maintains the community at a low level of complexity. All measured biological components are consistent with sedimentological and radiochemical studies which indicate a thin layer (<1cm) of biological mixing.

Benthic Sampling

Benthic samples were taken during four ten-day cruises aboard the R.V. *Planet* and R.V. *Helmsand*, German naval research ships, during spring 1993 and summer 1994. Benthic sampling was focused around the acoustic tower deployment sites - Old Tower, New Tower - in the central basin (Fig. 5). Other samples were taken from the landward and seaward ends of the bay. These included both muddy sites - Eckernfoerde Navy Base, Stanic Tower, Hausgarten - and one sandy site, Mittelgrund. Three or more replicate box cores (20 x 30 x 50 cm) were collected per station. Three subcores were taken from each box core, were extruded, and divided into 0-2 and 2-10 cm sections. Animals were seived from sediment using a 500-µm sieve and preserved. Animals were identified to the lowest possible taxonomic level (usually species). Each set of organisms was assigned to one of the following functional groups: surface deposit feeder, head-down deposit feeder, suspension feeder, or carnivore (see Table 4).

A bioturbation experiment was conducted to compare vertical particle mixing rates in different sediments and benthic communities. Three 15-cm diameter cores were collected from four stations along the axis of the bay - New Tower, Old Tower, Eckernfoerde Navy Base, Mittelgrund - in April 1993. Cores were maintained in a running seawater sytem at a shore lab, and subsampled as discussed previously. Samples were analyzed for fluorescence of tracer particles (modified from Carey, 1989). All values were corrected for background fluorescence.

Values from each depth interval were expressed as percent of the total fluorescence in that subcore.

Table 4. Rank order of dominant macrofauna in Eckernfoerde Bay.

| Polydora ciliata | polychaete | surface deposit feeder/facultative suspension feeder |
|-------------------------|-------------|--|
| Abra alba | bivalve | surface deposit feeder/facultative suspension feeder |
| Diastylus rathkei | crustacean | surface deposit feeder |
| Capitella sp. | polychaete | small head-down deposit feeder |
| Heteromastus filiformis | polychaete | small head-down deposit feeder |
| Pectinaria koreni | polychaete | large head-down deposit feeder |
| tubificid sp. | oligochaete | large head-down deposit feeder |
| Scoloplos armiger | polychaete | large head-down deposit feeder |
| Anaitides maculata | polychaete | carnivore |
| syllid sp. | polychaete | carnivore |
| Nepthys sp. | polychaete | carnivore |
| Harmathoe sp. | polychaete | carnivore |
| Sigambra sp. | polychaete | carnivore?scavenger |
| Mytilus edulis | bivalve | epifaunal suspension feeder |
| Cerastoderma sp. | bivalve | infaunal suspension feeder |
| Corbula gibba | bivalve | infaunal suspension feeder |

Biological Observations

Macrofaunal abundance was high throughout the bay, dominated by small adults and juveniles of larger species. The dominant macrofauna consisted of 16 species (90% of total species) and 4 functional groups (Table 4). Benthic samples from the muddy stations - Old Tower, New Tower - in both 1993 and 1994 were dominated by small, numerous surface deposit feeders. The polychaete *Polydora ciliata* and the tellinid bivalve *Abra alba* accounted for 75-90% of all animals in many samples. Most of the *A. alba* were juveniles (<2.5 mm). Adult *Abra alba*

were absent but their characteristic fecal pellets were abundant in the central basin; their presence indicates lateral transport from the sides of the basin. The surface deposit feeding cumacean, *Diastylus rathkei*, was numerically important in the June 1994 samples.

Abundances of surface deposit feeders in March-May, 1993 samples (27,000 - 73,000 m⁻²) were significantly greater than the June-July, 1994 abundances (1,700 - 9,000 m⁻²) (Fig. 6). Decreases in abundance were coupled with a shift in community composition at the muddy stations (Tower stations, Navy Base, Transect Station A, Hausgarten). The relative abundance of surface deposit feeders at the tower stations decreased from 80-90% in March-May 1993 to 60-85% in June 1994 (Fig. 7). Carnivores, especially *Harmathoe* sp. were proportionally more important in the June 1994 samples. The relative abundances of head-down deposit feeders remained unchanged between the two sampling years.

Mittelgrund is a sandy site near the mouth of Eckernfoerde Bay (see Fig. 6). This area supported a distinctly different fauna with significantly lower abundances than the muddy stations in the central basin. Mittelgrund had more head-down deposit feeders and a higher proportion of carnivores than the Tower sites (Fig. 7). Suspension feeders were also significantly more abundant in the sandier site.

The results of the particle mixing experiments showed that fluorescent particles were mixed approximately 0.5-1.0 cm into the sediment for all sites over a two-week period (Fig. 8), regardless of functional group.

Our data indicates that the benthic fauna of Eckernfoerde Bay is dominated by small, surface deposit feeders, which do not mix particles deeply but probably mix the top cm of the sediment on short time scales (days). Differences in mean abundances between the two sampling years is thought to be seasonal (March-May in 1993 versus late June in 1994) and is consistent with previous work done in the Kiel Bight (e.g. Bosselman 1988, Meyer-Reil and others 1987, Rumohr and Arntz 1982).

The abundances and composition of benthic fauna in Kiel Bight is strongly linked to seasonal patterns of oxgen availability in bottom waters (Kolmel 1979; 1977; Reimers; 1976). The benthic fauna of Kiel Bight typically experience an exponential increase in abundance, dominated by small, tube-building polychaetes such as *Polydora ciliata*, from March to early June followed by a crash in June-July (Weigelt 1991; Bosselman 1988; Meyer-Reil and others 1987;

Rumohr and Arntz 1982). In Eckernfoerde Bay, mean abundance of *P. ciliata* decreased an order of magnitude from the March-May 1993 sampling to the late June 1994 samples.

The primary controls on the benthic fauna of Eckernfoerde Bay during the study period were thought to be the rate of organic input to sediments, water-column stratification, deep-water mixing, and oxygen availability. During the spring bloom, carbon transport to the seafloor is between 10 and 15 g C m⁻² and one-third of the yearly organic input can reach the seafloor of the Kiel Bight in 1-2 weeks (Meyer-Reil and others 1987). Portions of the Kiel Bight, particularly the southwest portion where Eckernfoerde Bay is located, are annually subjected to periods of hypoxia and anoxia in summer due to stratification of the water column and stagnant physical conditions (Weigelt 1991; 1990; Meyer-Reil and others 1987). We hypothesize that the low oxygen conditions that occur with water column stratification in Eckernfoerde Bay causes a regular disturbance which controls the complexity of benthic community structure.

All measured components of Eckernfoerde Bay - abundance, diversity, animal size, functional groups, particle mixing - point to a community dominated by pioneering species (sensu Rhoads and Boyer 1982), especially *Polydora ciliata* and *Capitella* sp., and a system controlled by a regular or recent disturbance that reduced the complexity of the community structure and sediment reworking by the benthic fauna. This is borne out by our bioturbation experiments (Fig. 8), our sedimentology and radiochemistry results, and with numerous German studies of the Kiel Bight (e.g. Weigelt 1991, Bosselman 1988, Meyer-Reil and others 1987, Rumohr and Arntz 1982).

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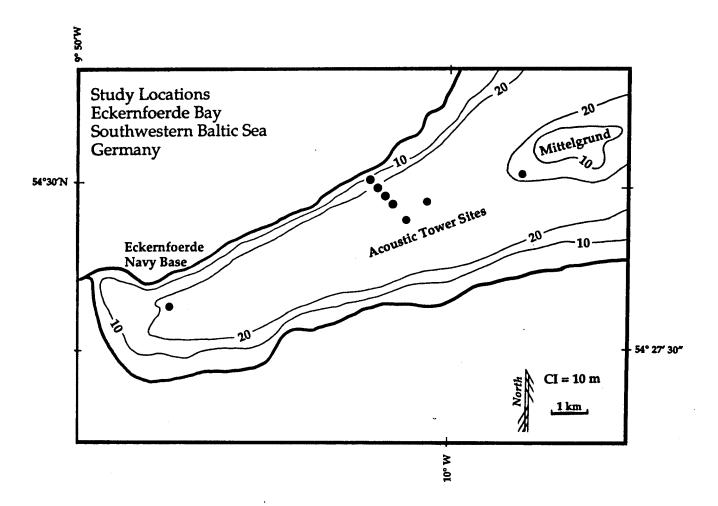


Figure 1 Map of coring locations in Eckernfoerde Bay.

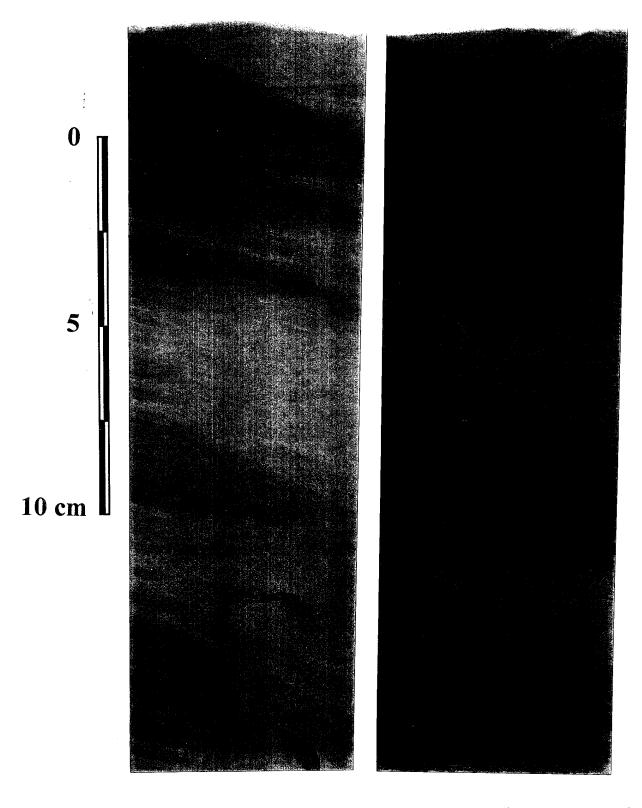


Figure 2 X-radiograph positives, BS4-611 (A) and BS4-602 (B). Stations were separated by ~1 km. Note darker laminations (non-pelletized event layers). Thicker, light-shaded beds are intensely pelletized.

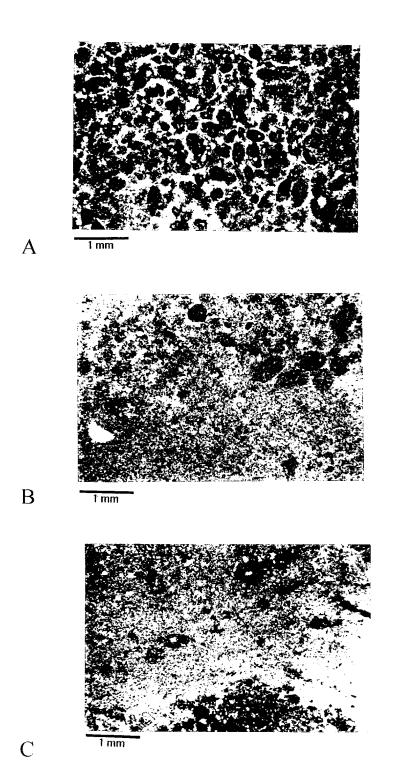
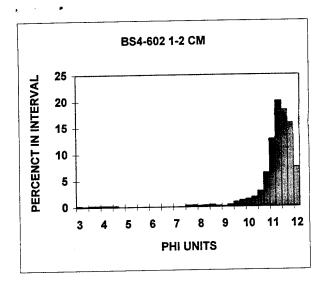


Figure 3 Photomicrographs, BS4-602. Field of view is 5.5 mm wide. A) 2.5-2.8 cm depth, plane light: intensely pelletized horizon; fecal pellets produced by the surface-deposit-feeding bivalve *A. alba* B); 6.7-7.0 cm depth, crossed polars: upper portion of event layer showing transition from non-pelletized (lower) to pelletized fabric (upper); C) 7.0-7.3 cm depth, crossed polars: basal contact between non-pelletized event layer and underlying pelletized horizon



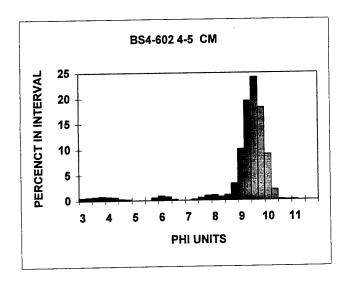


Figure 4 Grain size distributions BS4-602, 1-2 cm; and BS4-602, 3-4 cm. The 1-2 cm interval is characteristic of pelletized intervals, and the 3-4 cm interval, slightly coarser, is characteristic of non-pelletized event layers.

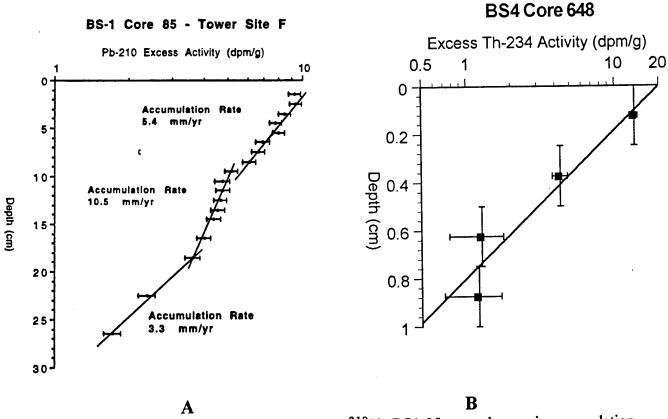


Figure 5 Radiochemical profiles. A)Excess ²¹⁰Pb, BS1-85, note changes in accumulation rate through time; B) excess ²³⁴Th, BS4-648, excess activities are restricted to the upper 9 mm.

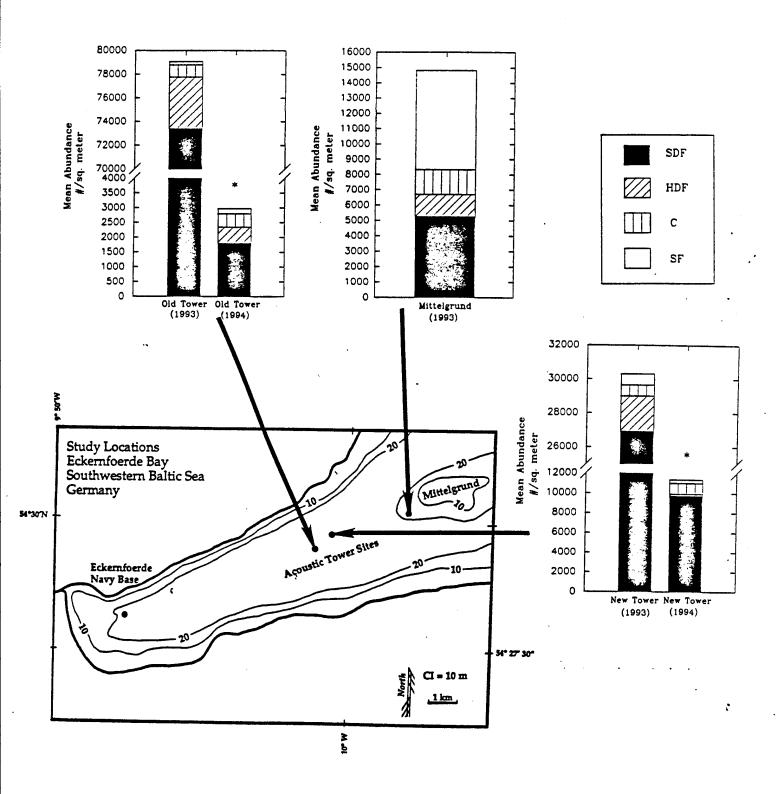


Figure 6. Mean Abundances at Primary Study Sites, 1993: mean from three cruises in Spring, 1993, 1994: mean from one cruise in Summer, 1994. SDF = surface deposit feeders; HDF = head-down deposit feeders; C = carnivores; SF = suspension feeders; B = browsers. * = significantly different abundances than 1993 samples (α=0.05).

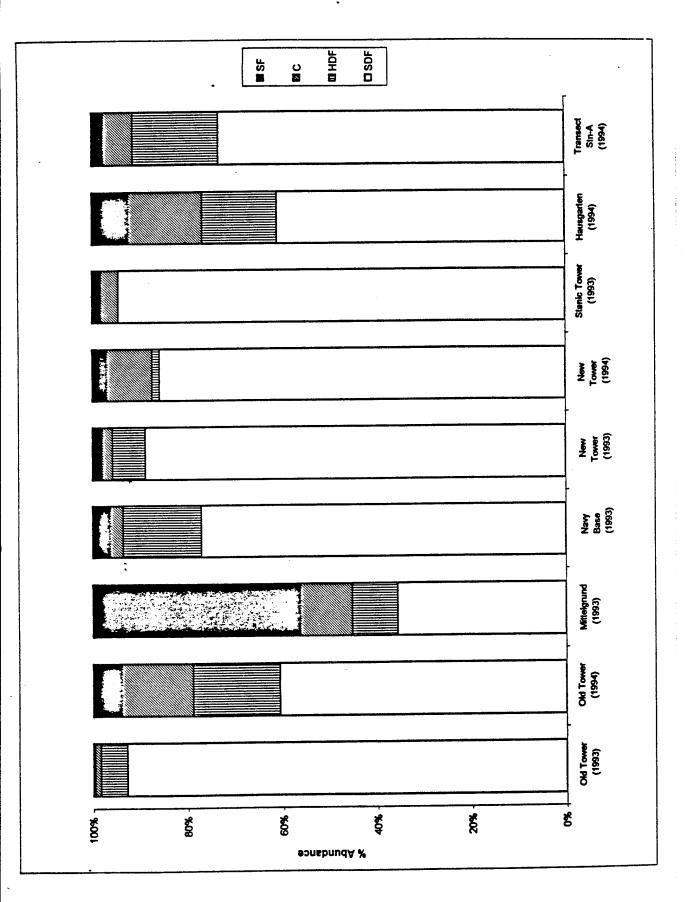


Figure 7. Percent Composition of Functional Groups by Station. SDF = surface deposit feeders; HDF = head-down deposit feeders; C = carnivores; SF = suspension feeders; B = browsers.

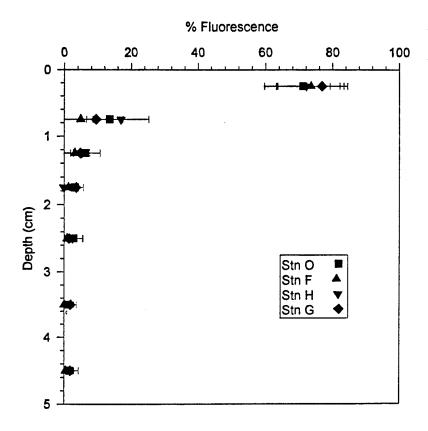


Figure 8. Vertical profiles of percent fluorescence (means and standard deviations) at primary study sites. Values are corrected for background fluorescence. Stations: O = New Tower, F = Old Tower, H = Eckernfoerde Navy Base, G = Mittelgrund.

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